

### **Amendments to the Claims:**

This listing of claims will replace all prior versions, and listings, of claims in this application.

### **Listing of Claims:**

1. (Currently Amended) A composition comprising a therapeutically effective amount of an antisense molecule targeted to the sequence set forth in SEQ ID NO:1 chemically linked to one or more moieties that enhance cellular uptake of said antisense molecule in to a cell admixed with an acceptable carrier.

2. (Currently Amended) A composition comprising a therapeutically effective amount of an antisense molecule targeted to the sequence set forth in SEQ ID NO:2 chemically linked to one or more moieties that enhance cellular uptake of said antisense molecule in to a cell admixed with an acceptable carrier.

3. (Currently Amended) A composition comprising a therapeutically effective amount of an antisense molecule comprising the nucleotide sequence set forth in SEQ ID NO:3, said sequence complementary to nucleotides 156-185 of BC200 RNA, and is chemically linked to one or more moieties that enhance cellular uptake of said antisense molecule in to a cell admixed with an acceptable carrier.

4. (Currently Amended) A composition comprising a therapeutically effective amount of an antisense molecule comprising the nucleotide sequence set forth in SEQ

ID NO:4, said sequence complementary to nucleotides 158-178 of BC200 RNA, and is chemically linked to one or more moieties that enhance cellular uptake of said antisense molecule in to a cell admixed with a-an acceptable carrier.

5. (Withdrawn) An isolated antisense molecule comprising the nucleotide sequence set forth in SEQ ID NO:5.

6. (Withdrawn) An isolated nucleic acid molecule comprising the nucleotide sequence set forth in SEQ ID NO:6, complementary to DNA encoding BC200 RNA.

7. (Cancelled).

8. (Withdrawn) A method for treating a neurological disorder or cancer in a subject, said method comprising down-regulating BC200 RNA in the subject.

9. (Withdrawn) The method of claim 8 wherein the down-regulating of BC200 RNA in a subject comprises administering a therapeutically effective amount of a dominant negative mutant of BC200 RNA or a small interfering RNA.

10. (Withdrawn) The method of claim 8 wherein the down-regulating of BC200 comprises administering a therapeutically effective amount of an antisense molecule targeted to the nucleotide sequence set forth in SEQ ID NO:1 or SEQ ID NO:2.

11. (Withdrawn) The method of claim 8 wherein the down-regulating of BC200 comprises administering a therapeutically effective amount of at least one of SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, or SEQ ID NO:6.

12. (Withdrawn) The method of any one claims 8-11 wherein the neurological disorder is at least one of Alzheimer's disease, Fragile X mental retardation syndrome, Down's syndrome and Parkinson's disease.

13. (Withdrawn) The method of any one of claims 8-11 wherein the cancer is at least one of squamous cell carcinoma of the tongue and lung, epithelial carcinoma of the esophagus, tubular adenocarcinoma of the stomach, breast adenocarcinoma, adenocarcinoma of the lung, mucoepidermoid of the parotid gland, melanoma of the skin, papillary carcinoma of the ovaries, or endothelial adenocarcinoma of the cervix.

14. (Withdrawn) A method for treating epilepsy in a subject, the method comprising up-regulating BC200 RNA in a patient.

15. (Withdrawn) The method of claim 14 wherein the up-regulating comprises administering to the patient a therapeutically effective amount of BC200 RNA.

16. (Withdrawn) The method of claim 14 wherein the up-regulating comprises administering to the patient a gene therapy construct having a DNA or RNA

corresponding to BC200 operably linked to a promoter which functions in the cells of the subject.

17. (Currently Amended) A kit comprising a therapeutically effective amount of an antisense molecule of any one of claims 1-4 chemically linked to one or more moieties that enhance cellular uptake of said antisense molecule in to a cell and an acceptable carrier, wherein the antisense molecule is admixed with the carrier.

18. (Currently Amended) The kit of claim 17 wherein the therapeutically effective amount of the antisense molecule is packaged separately from the acceptable carrier.

19. (Cancelled)

20. (Cancelled)